AMENDMENT US APPLN. NO. 10/626,138

## Amendments to claims:

## Listing of Claims as amended:

1. (Currently Amended) A method for the treatment or prevention of heart failure, which comprises the administration to a patient in need thereof of a pharmaceutical composition comprising cilobradine or a pharmaceutically acceptable salt thereof, together with a pharmaceutically suitable carrier.

## 2-3 Cancelled.

- 4. (Original) The method of claim 1 wherein the galenical formulation of the pharmaceutical composition is a tablet, a drinking solution, a capsule, a suppository or an injectable formulation.
- 5. (Original) The method of claim 4 wherein the galenical formulation of the pharmaceutical composition is a tablet.
- 6. (Original) The method of claim 4 wherein the galenical formulation of the pharmaceutical composition is a drinking solution.
- 7. (Currently Amended) The method of claim 1 wherein the pharmaceutical composition is administered following a single daily application scheme.
- 8. (Original) The method of claim 7 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.01 and 20 mg/kg body weight.

- 9. (Original) The method of claim 7 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.05 and 5 mg/kg body weight.
- 10. (Original) The method of claim 7 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.1 and 2.5 mg/kg body weight.
- 11. (Original) The method according to claim 7 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.1 and 1 mg/kg body weight.
- 12. (Original) The method of claim 7 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.1 and 0.75 mg/kg body weight.
- 13. (Currently amended) The method of claim 1 wherein the treatment or prevention of heart failure is performed in combination with other therapeutic agents for the treatment or prevention of heart failure.
- 14. (Original) The method of claim 13 wherein said therapuneutic agents include diuretics, cardiac glycosides, ACE inhibitors, ARBs, vasodilators, beta blockers or inotropes.
- 15. (New) The method of claim 1 wherein the pharmaceutical composition is administered multiple times daily.
- 16. (New) The method of claim 15 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.01 and 20 mg/kg body weight.
- 17. (New) The method of claim 15 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.05 and 5 mg/kg body weight.

AMENDMENT US APPLN. NO. 10/626,138

- 18. (New) The method of claim 15 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.1 and 2.5 mg/kg body weight.
- 19. (New) The method according to claim 15 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.1 and 1 mg/kg body weight.
- 20. (New) The method of claim 15 wherein the administered dose of cilobradine or its pharmaceutically acceptable salt is between 0.1 and 0.75 mg/kg body weight.